

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

In re Application of Alan SOLOMON *et al.* Confirmation No.: 7724  
Serial No.: 09/316,387 Group Art Unit: 1649  
Filed: May 21, 1999 Examiner: EMCH, Gregory  
For: METHODS FOR AMYLOID REMOVAL USING ANTI-AMYLOID ANTIBODIES

**DECLARATION UNDER 37 C.F.R. § 1.132**

I, Mr. Stephen Wood, hereby declare as follows:

1. I have been employed by Amgen and I am presently a Senior Scientist at Amgen. During my Employment at Amgen, I have been engaged in research and development in the area of amyloid related diseases, including Alzheimer's disease. I have a M.S. in Biochemistry from The University of Nebraska-Lincoln. My CV is attached as Exhibit 5.

2. The following experiments were performed at Amgen, Inc. under my supervision:

3. Binding of mAb 369.2B (the antibody disclosed the cited document WO 96/25435) to A $\beta$  was compared to control antibody 2.1 chimera (disclosed as "PC2" in US 2003/0082191). In these experiments, A $\beta$  1-40 (A $\beta$ 40) fibril or monomer, A $\beta$  1-42 (A $\beta$ 42) fibril or monomer,  $\alpha$ -synuclein or collagen fibrils were coated onto 96-well microtiter plates and dried overnight at 37°C. Then, mAb 369.2B or mAb 2.1 chimera were incubated at various dilutions in the plates. Binding of the antibodies was detected using a biotinylated secondary antibody followed by europium-tagged streptavidin. The assay readout is the time-resolved fluorescence of europium (TRF). These results are illustrated on Exhibits 1 and 2.

4. Exhibits 1C and D are control experiments that demonstrate that 369.2B and 2.1 chimera antibodies do not exhibit non-specific binding. Exhibits 1A and Exhibit 2A depict experiments showing that mAb 369.2B does not bind to A $\beta$  1-40 (A $\beta$ 40) fibrils or monomers, while 2.1 chimera does.

5. Exhibits 1B and 2B show binding of said antibodies to A $\beta$  1-42 fibrils and monomers. These data show that mAb 369.2B exhibits weaker binding compared to 2.1 chimera, as evidenced by a shift in the curve of mAb 369.2 to the right.

6. Immunohistochemistry was performed to demonstrate whether mAb 369.2B bound to amyloid plaques in unfixed brain tissue. Fresh-frozen sections from an 18 month-old Tg2576 mouse, not fixed with fixative or treated with formic acid, were incubated with 1mg/ml of the indicated antibodies. Binding was detected using biotinylated goat anti-mouse IgG and avidin-HRP with Mouse to mouse system (Vector). Images were taken at 4x magnification on a Nikon Microphot-FXA microscope. A 6-score system was assigned based on staining level by visual assessment, where 0 represents the staining level seen in corresponding control sections and 5 plus are densest staining. mAb 4G8 binds to the mid-region of A $\beta$  (residues 17-24), mAb 369.2b is a C-terminal A $\beta$  binder, mAb 2.1 is a N-terminal binder and mAb 4D2 is an irrelevant IgG1 (negative control). These data are depicted on Exhibit 3. These data show that there is little binding of mAb 369.2B to plaques in the brain tissue, when compared to mAb 2.1. These data confirm the results shown in Exhibits 1B and 2B, that mAb 369.2B has low binding affinity for amyloid fibrils. The low-level binding exhibited by mAb 369.2b is not sufficient to opsonize amyloid fibrils and induce removal of amyloid deposits *in vivo*.

7. *Ex vivo* phagocytosis assay was used as an efficacy test for the antibodies. This assay translates biochemical-binding properties into therapeutic utility by evaluating an antibody's capacity to induce phagocytosis of amyloid plaques. It has been shown previously that the

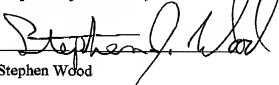
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phagocytosis assay is the best available predictor of *in vivo* antibody efficacy (reduction of plaque burden) in mouse models of Alzheimer's Disease.<sup>1</sup>

8. Cell binding and tissue binding assays were performed by increasing amounts of the indicated antibodies incubated with unfixed brain sections from a 19 month-old Tg2576 mouse and microglial IC21 cells (a phagocytic cell-line). After 24-hour incubation, the tissue samples were fixed, and stained with biotinylated 6E10 followed by a streptavidin-FITC (Vector). IC-21 cells were stained with rat anti-CD45 and goat anti-rat IgG-cy3. Blinded slides were scored for phagocytic events on a Zeiss LSM5 Pascal Confocal Microscope and reported as percent of total plaques counted (over 100) that were partially or fully engulfed ( $n=3$ ). As shown in Exhibit 3, increasing the concentration of mAb 2.1 resulted in a concentration-dependent increase in phagocytosis with  $EC_{50}$  value, 0.68  $\mu\text{g/ml}$ . Phagocytic efficacies of  $A\beta$  mid-region binder, mAb 4G8 and C-terminal  $A\beta$  binders, mAb 369.2b and Confab 42, were substantially reduced when compared to the N-terminal mAb 2.1. The minimal level of phagocytic activity induced by mAb 369.2b would not be predicted to remove amyloid deposits *in vivo*.

9. I declare that all statements made herein on my own knowledge are true and that all statements made on information and belief are believed to be true and further that these statements are made with the knowledge that willful false statements and the like are punishable by fine or imprisonment, or both, under § 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the above-referenced application or any patent issuing thereon.

Respectfully submitted,

  
Stephen Wood

Date:

8/6/08

<sup>1</sup> Bard, F. *et al.* (2003) PNAS 100(4), 2023-8 (Exhibit 6).

EXHIBIT 1

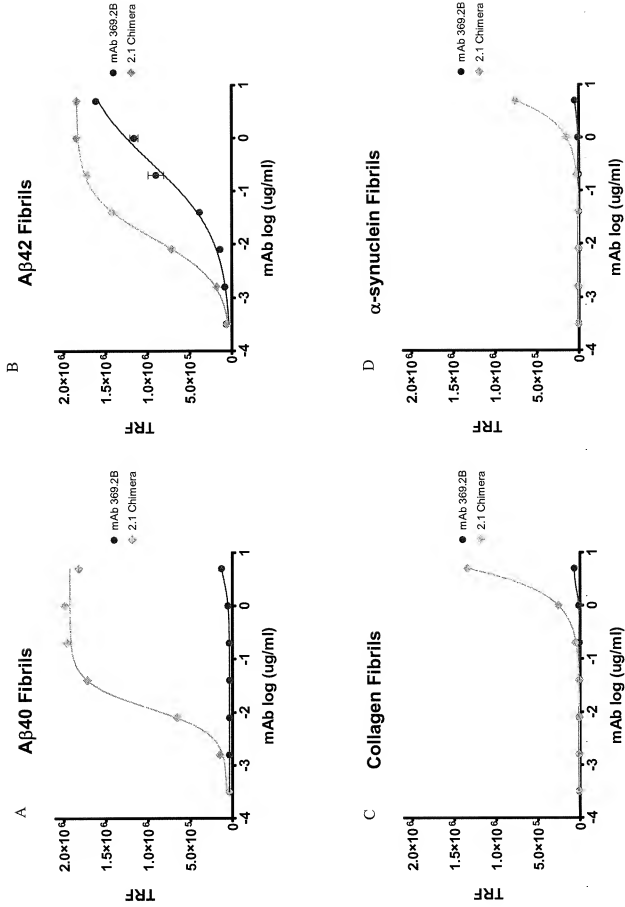
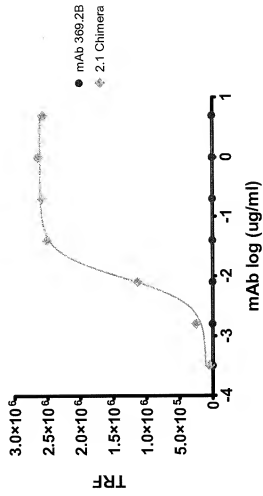


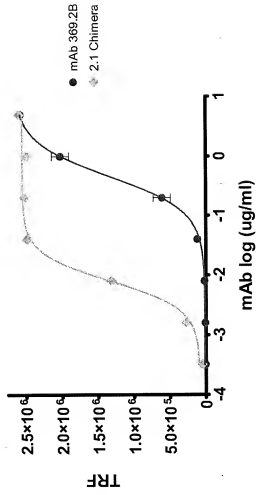
EXHIBIT 2

**A $\beta$ 40 Monomer**



A

**A $\beta$ 42 Monomer**



B

Exhibit 3

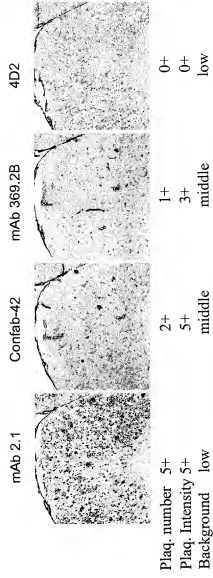
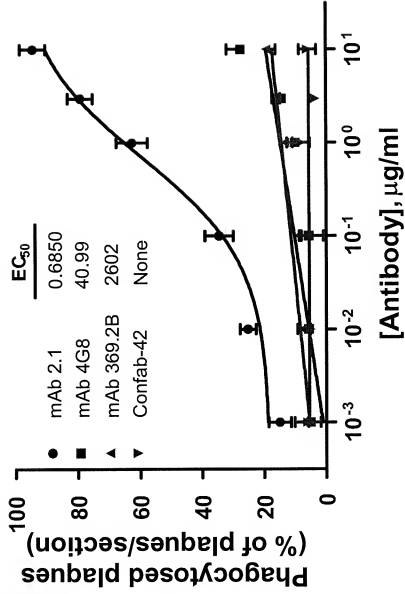


EXHIBIT 4



**Stephen J. Wood****EDUCATION:**

Bachelor of Science in Biological Sciences (1987), University of Nebraska-Lincoln  
Master of Science in Biochemistry (1990), University of Nebraska-Lincoln

**Professional Experience:**

|                |  |
|----------------|--|
| 9/04 – Present | Senior Scientist (formerly Research Scientist II), Neuroscience Dept., Amgen Thousand Oaks, CA                     |
| 9/02 – 9/04    | Research Scientist I, Neuroscience Dept., Amgen Thousand Oaks, CA  |
| 2/01 – 2/02:   | Associate Scientist II, Neuroscience Dept., Amgen Thousand Oaks, CA  |
| 6/99 – 2/01:   | Associate Scientist I, Neuroscience Dept., Amgen Thousand Oaks, CA.  |
| 6/97 – 6/99:   | Research Associate III, Neuroscience Dept., Amgen Thousand Oaks, CA.   |
| 1994 - 6/97:   | Senior Scientist, Macromolecular Sciences Dept., SmithKline Beecham Pharmaceuticals, King of Prussia, PA 19406.    |
| 1992 - 1994:   | Scientist, Macromolecular Sciences Dept., SmithKline Beecham Pharmaceuticals, King of Prussia, PA 19406.           |
| 1990 - 1992:   | Associate Scientist, Macromolecular Sciences Dept., SmithKline Beecham Pharmaceuticals, King of Prussia, PA 19406. |

**Scientific Expertise:**

- Alzheimer's and Parkinson's Disease research
- Protein aggregation/amyloid fibril formation
- *In vitro* assay development and optimization
- Protein purification, modification and characterization
- Protein-protein interactions
- Protein mutagenesis studies
- Protein therapeutic candidate generation
  - Primary screening (phage and yeast display; antibody generation)
  - Candidate evaluation (specificity and affinity measurements, epitope mapping, etc.)
- Small molecule therapeutic candidate generation



- o Primary, high-throughput assay development and screening
- o Candidate evaluation (secondary screens, IC<sub>50</sub> measurements, etc.)

## **PUBLICATIONS:**

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3. Myszka, D. G., Wood, S. J. and Biere, A. L. (1999). Analysis of Fibril Elongation Using Surface Plasmon Resonance Biosensors. *Methods in Enzymology* 309 (25), pp. 386-402
4. Wood, S. J., Wypych, J., Steavenson, S., Louis, J. C., Citron, M. and Biere, A. L. (1999).  $\alpha$ -Synuclein Fibrillogenesis is Nucleation-Dependent: Implications for the Pathogenesis of Parkinson's Disease. *Communication in Journal of Biological Chemistry* 274 (28), pp 19509-19512.
5. Howlett, D. R., Perry, A. E., Godfrey, F., Swatton, J. E., Jennings, K. H., Spitzfaden, C., Wadsworth, H., Wood, S. J. and Markwell, R. E. (1999). Inhibition of Fibril Formation in  $\beta$ -Amyloid Peptide by a Novel Series of Benzofurans. *Biochemical Journal* 340, pp 283-289.
6. Narhi, L.\*, Wood, S. J.\*, Steavenson, S.\*, Yijia, J., Gay, M. U., Anafi, D., Kaufman, S. A., Martin, F. Sitney, K., Denis, P., Louis, J. C., Wypych, J., Biere, A. L., Citron, M. (1999). Both Familial Parkinson's Disease Mutations Accelerate  $\alpha$ -Synuclein Aggregation. *Journal of Biological Chemistry* 274 (14), pp 9843-9846.
7. Wood, S.J., Chan, W. and Wetzel, R. (1996). An ApoE-A $\beta$  Inhibition Complex of A $\beta$  Fibril Extension. *Chemistry and Biology* 3 (11), pp 949-956.
8. Wood, S.J., Chan, W. and Wetzel, R. (1996). Fibril Formation Induced by Exogenous Beta-A4 Seeds is Inhibited by Apolipoprotein E. *Biochemistry* 35 (38), pp 12623-12628.
9. Wood, S.J., MacKenzie-LoCastro, L., Holl, W., Maleeff, B., Hurle, M.R. and Wetzel, R. (1996). Selective Inhibition of Beta-A4 Fibril Formation. *Journal of Biological Chemistry* 271 (8), pp 4086-4092.
10. Wood, S.J., Maleeff, B., Hart, T. and Wetzel, R. (1996). Physical, Morphologic and Functional Differences Between pH 5.8 and 7.4 Aggregates of the Alzheimer's Peptide A-Beta. *Journal of Molecular Biology* 256, pp 870-877.

11. Wood, S.J., Wetzel, R. and Hurle, M.R. (1995). Proline Substitutions Eliminate the Amyloidogenicity of Peptides Analogous to the 12-26 Region of Beta-A4. *Biochemistry* 34 (3), pp 724-730.
12. Howlett, D.R., Jennings, K.H., Lee, D.C. Clark, M.S.G., Brown, F., Wetzel, R.B., Wood, S.J., Camilleri, P., and Roberts, G.W. (1995). Aggregation State and Neurotoxic Properties of Alzheimer Beta-Amyloid Peptide. *Neurodegeneration* 4, pp 23-32.
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## **ABSTRACTS**

1. Wood, S.J., Jaronsinski, M., and Biere, A.L. Effects of Single Amino Acid Substitutions on A $\beta$  Fibril Elongation. FASEB Conference on Amyloids and Other Protein Assembly Processes, Copper Mountain, CO, June 11-16, 2000.
2. Wood, S.J., Wetzel, R.B. and Hurle, M.R. Effects of Mutation and pH on Fibrillization of Truncated Beta-A4 Peptides. The VII<sup>th</sup> International Symposium on Amyloidosis, Kingston Ontario Canada, July 11-15, 1993. Fibrillogenesis #W01, pp 156.

# Epitope and isotype specificities of antibodies to $\beta$ -amyloid peptide for protection against Alzheimer's disease-like neuropathology

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Elan Pharmaceuticals, 800 Gateway Boulevard, South San Francisco, CA 94080

Edited by L. L. Iversen, University of Oxford, Oxford, United Kingdom, and approved December 18, 2002 (received for review October 16, 2002)

**Transgenic PDAPP mice, which express a disease-linked isoform of the human amyloid precursor protein, exhibit CNS pathology that is similar to Alzheimer's disease. In an age-dependent fashion, the mice develop plaques containing  $\beta$ -amyloid peptide (A $\beta$ ) and exhibit neuronal dystrophy and synaptic loss. It has been shown in previous studies that pathology can be prevented and even reversed by immunization of the mice with the A $\beta$  peptide. Similar protection could be achieved by passive administration of some but not all monoclonal antibodies against A $\beta$ . In the current studies we sought to define the optimal antibody response for reducing neuropathology. Immune sera with reactivity against different A $\beta$  epitopes and monoclonal antibodies with different isotypes were examined for efficacy both *ex vivo* and *in vivo*. The studies showed that: (i) of the purified or elicited antibodies tested, only antibodies against the N-terminal regions of A $\beta$  were able to invoke plaque clearance; (ii) plaque binding correlated with a clearance response and neuronal protection, whereas the ability of antibodies to capture soluble A $\beta$  was not necessarily correlated with efficacy; (iii) the isotype of the antibody dramatically influenced the degree of plaque clearance and neuronal protection; (iv) high affinity of the antibody for Fc receptors on microglial cells seemed more important than high affinity for A $\beta$  itself; and (v) complement activation was not required for plaque clearance. These results indicate that antibody Fc-mediated plaque clearance is a highly efficient and effective process for protection against neuropathology in an animal model of Alzheimer's disease.**

Immunization of the transgenic PDAPP mice with  $\beta$ -amyloid peptide (A $\beta$ )-derived immunogens results in an antibody response that facilitates the clearance of plaques within the central nervous system (CNS) (1–4). Although a number of mechanisms are likely to operate in this clearance response (5, 6), our previous findings strongly indicate that antibody-mediated, Fc-dependent phagocytosis by microglial cells and/or macrophages is important to the process (7). Importantly, a T cell response was not required for amyloid plaque clearance. When peripherally administered, antibodies against A $\beta$  entered the CNS of PDAPP transgenic mice, decorated amyloid plaques, and induced plaque clearance. Comparing different antibodies in an *ex vivo* assay with sections of PDAPP or Alzheimer's disease (AD) brain, there was a strong correlation between those that produced *ex vivo* efficacy and those that were efficacious *in vivo*. Fc receptors on microglial cells were found to be key for the clearance response in this assay. However, it has been reported that antibody efficacy can also be obtained *in vivo* by mechanisms that are independent of Fc interactions (8). Studies have indicated that an antibody directed against the midportion of A $\beta$ , which cannot recognize amyloid plaques, appears to bind to soluble A $\beta$  and reduce plaque deposition (6). In addition, it has been reported recently that short-term treatment with this antibody improved performance in an object-recognition task without affecting amyloid burden (9).

To understand the parameters of an antibody response that are required for neuronal protection, several questions should be considered. Is neuronal protection associated with plaque clearance, or is it necessary for antibodies to capture soluble aggregates of A $\beta$  to protect neurons against the directly toxic effects of the peptide? Does a clearance response depend on Fc receptor-mediated phagocytosis of A $\beta$  after antibody binding or on complement receptor-mediated phagocytosis after antibody binding and complement activation? Alternatively, is a clearance response independent of antibody Fc receptor function?

In the current study we approached these questions by examining the influence of different antibody epitopes and isotypes on plaque clearance and neuronal protection. The studies took advantage of the fact that some epitopes of A $\beta$  are preferentially available for antibody binding within plaques, whereas others are only available for antibody capture of the soluble peptide. In addition, the isotype of an antibody is important for either Fc or complement-mediated phagocytosis of A $\beta$  by microglial cells, because antibody isotype defines its affinity for Fc receptors as well as its ability to activate complement. If plaque clearance and/or neuronal protection do not depend on Fc-mediated processes, then the isotype of an antibody against A $\beta$  should have little impact on efficacy. These studies provide insight for the design of antibodies with therapeutic potential.

## Materials and Methods

**A $\beta$  Fragments.** Peptides corresponding to A $\beta$ 1–5, A $\beta$ 3–9, A $\beta$ 5–11, and A $\beta$ 15–24 and the reverse sequence A $\beta$ 5–1 were synthesized contiguous to a 17-aa T cell epitope derived from ovalbumin (amino acids 323–339, ISQAVHAAHAEINEAGR) on a branched peptide framework (triple-lysine core with four peptide arms) to produce a multiantigen peptide as described (10). Polyclonal antibodies against A $\beta$ 1–42 (pAb 1–42) were raised and the Ig fraction was isolated as described (7). pAb-EL16, pAb-EL17, and pAb-EL20 were obtained from the sera of PDAPP mice immunized with peptides corresponding to A $\beta$ 1–7, A $\beta$ 15–24, and A $\beta$ 3–9, respectively, which had been synthesized on a branched framework as described above. pAb-EL26 was obtained from the sera of mice immunized with A $\beta$ (7–1)–42. The peptides were synthesized by AnaSpec (San Jose, CA).

**Monoclonal Antibodies (mAbs).** The production of mAbs 10D5 and 6C6, which were raised against synthetic A $\beta$ 1–28 coupled to a carrier protein, has been described (11). mAbs 12B4, 2C1, 12A11, and 3A3 were raised against synthetic A $\beta$ 1–42 by using similar methodology except that hybridoma supernatants were screened by an RIA. All antibodies were purified by HPLC and were free of endotoxin (<1 endotoxin unit/mg protein) as

This paper was submitted directly (Track II) to the PNAS office.

Abbreviations: AD, Alzheimer's disease; A $\beta$ ,  $\beta$ -amyloid peptide; pAb, polyclonal Ab.

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determined by the *Limulus* amoebocyte gel-clot assay (Associates of Cape Cod). The mAbs 3D6 and 21F12 were obtained as described (7), and 22D12 and 266 were raised against synthetic A $\beta$ 13–28 (12).

**Epitope Mapping.** Epitope mapping of the mAbs and pAbs was performed by using an ELISA that measured antibody binding to overlapping peptides (10 amino acid peptides offset by 1 residue) covering the entire A $\beta$ 1–42 sequence. The first 32 peptides were biotinylated at the C terminus, and the last 10 peptides were biotinylated at the N terminus. The biotinylated peptides were synthesized by Mimotopes (Clayton, Victoria, Australia) and captured on streptavidin-coated wells of a 96-well plate (Pierce).

**Passive and Active Immunization Procedures.** mAbs in PBS were given via passive administration (i.p. injection) at a dose of 10  $\mu$ g/kg weekly for 6 months. For active immunization, 100  $\mu$ g of A $\beta$  fragment was administered by i.p. injection in complete Freund's adjuvant followed by boosts with 100  $\mu$ g of peptide in incomplete Freund's adjuvant at 2 and 4 weeks, and monthly thereafter.

**Antibody Binding to Aggregated and Soluble A $\beta$ 1–42.** Serum titers (determined by serial dilution) and mAbs binding to aggregated synthetic A $\beta$ 1–42 were performed by ELISA as described (1). Soluble A $\beta$ 1–42 refers to the synthetic A $\beta$ 1–42 peptide sonicated in dimethyl sulfoxide. Serial dilutions of sera or mAb at 20  $\mu$ g/ml were incubated with 50,000 cpm [<sup>125</sup>I]A $\beta$ 1–42 (~190  $\mu$ Ci/ $\mu$ mol; labeling with Iodogen reagent, Pierce) overnight at room temperature. Fifty microliters of a slurry containing 75  $\mu$ g/ml protein A Sepharose (Amersham Pharmacia) and 200  $\mu$ g of rabbit anti-mouse IgG (H+L) (Jackson ImmunoResearch) was incubated with the diluted antibodies for 1 h at room temperature, washed twice, and counted on a Wallac gamma counter (Perkin-Elmer). All steps were performed in RIA buffer consisting of 10 mM Tris, 0.5 M NaCl, 1 mg/ml gelatin, and 0.5% Nonidet P-40, pH 8.0.

**Ex Vivo Assay.** Cryostat sections (10  $\mu$ m in thickness) of PDAPP mouse brain were thaw-mounted onto round polysilyne-coated coverslips and placed in the wells of 24-well tissue-culture plates. Microglial cells and antibodies were added to the wells and cultured for 24 h as described (7). After incubation, cultures were extracted with an 8 M urea buffer and frozen quickly. Total A $\beta$  level in the cultures was determined by ELISA as described (13).

Statistical analyses were performed by using PRISM 3.0 software (GraphPad, San Diego).

## Results and Discussion

**A $\beta$  Epitope: Epitopes Within the N Terminus of A $\beta$  Are Important for Plaque Clearance and Reduction of Neuritic Pathology.** We have shown previously that not all antibodies against A $\beta$  can trigger plaque clearance *in vivo*. Efficacy can be predicted by the ability of antibodies to both bind plaques within unfixed sections of PDAPP or AD brains and trigger plaque clearance in an *ex vivo* assay (7). In the current study, a number of mAbs and pAbs directed against different epitopes of A $\beta$  were examined for plaque reactivity and *ex vivo* efficacy. Only antibodies against epitopes within the N-terminal 11 aa of A $\beta$  were found to be active in either regard (Table 1). These findings also illustrate that, of the five antibodies previously examined *in vivo*, those against N-terminal epitopes were effective in reducing plaque burden (pAb1–42, 3D6, and 10D5), whereas those against C-terminal epitopes were inactive (16C11 and 21F12) (7).

To extend these findings and further characterize epitopes within the N terminus of A $\beta$ , a series of peptides were compared for their ability to trigger an efficacious antibody response *in vivo*. Twelve- to 13-month-old PDAPP mice were immunized

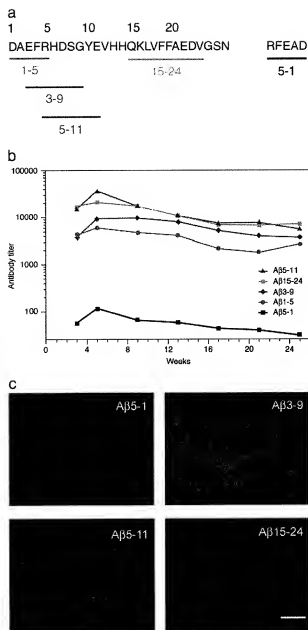
**Table 1. Antibodies directed against epitopes within the N-terminal 11 aa of A $\beta$ 1–42 bind amyloid plaques and trigger phagocytosis in an *ex vivo* assay**

| Antibody | Epitope | Binds plaques (PDAPP) | Triggers <i>ex vivo</i> phagocytosis |
|----------|---------|-----------------------|--------------------------------------|
| 3D6      | 1–5     | ++                    | ++                                   |
| pAb-EL16 | 1–7     | ++                    | ++                                   |
| pAb1–42  | 1–11    | ++                    | ++                                   |
| 10D5     | 3–7     | ++                    | ++                                   |
| pAb-EL21 | 5–11    | +                     | +                                    |
| pAb-EL26 | 11–26   | –                     | –                                    |
| 22D12    | 18–21   | –                     | –                                    |
| 266      | 16–24   | –                     | –                                    |
| pAb-EL17 | 15–24   | –                     | –                                    |
| 16C11    | 33–42   | –                     | –                                    |
| 21F12    | 34–42   | –                     | –                                    |

Both measures were scored on three-step visual rating system based on fluorescence intensity for the plaques and by degree of A $\beta$  uptake as described (7).

with one of three N-terminal peptide fragments (A $\beta$ 1–5, A $\beta$ 3–9, or A $\beta$ 5–11) or a fragment derived from an internal region of the peptide (A $\beta$ 15–24) (Fig. 1a). The internal peptide A $\beta$ 15–24 encompasses the epitope of antibody 266, which exhibits high affinity for soluble A $\beta$  (12), but as shown above it does not recognize plaques in sections of unfixed AD or PDAPP tissue (Table 1). Thus, it was of interest to determine whether a polyclonal response directed against this peptide could produce antibodies capable of plaque recognition or whether reactivity with soluble A $\beta$  alone was sufficient to provide efficacy. In these studies, a peptide with reverse sequence (A $\beta$ 5–1) served as a negative control. The peptides were synthesized conjugated to a 17-aa T cell epitope derived from ovalbumin and presented in an identical multivalent configuration (see *Materials and Methods*). All the peptides (except A $\beta$ 5–1 reverse mer) produced sera that recognized aggregated synthetic A $\beta$ 1–42 by ELISA (Fig. 1b). In contrast, only sera against the N-terminal peptides were able to recognize A $\beta$  within plaques; antisera against A $\beta$ 15–24 did not bind plaques despite strong reactivity with the synthetic aggregated peptide (Fig. 1c). There also were differences between the serum groups in their ability to capture soluble A $\beta$  (Fig. 2a). Less than 30% of the sera from mice immunized with A $\beta$ 1–5 or A $\beta$ 3–9 captured the soluble peptide (27% and 5%, respectively). In contrast, sera from approximately half of the animals immunized with A $\beta$ 5–11 and all of those immunized with A $\beta$ 15–24 captured soluble A $\beta$ 1–42.

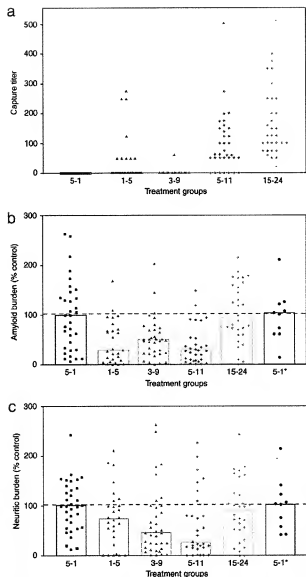
Because the degree of A $\beta$  deposition can vary greatly as PDAPP mice age, the *in vivo* study was designed with at least 30 animals per group. Efficacy data are shown for individual mice and expressed as the percentage of either amyloid burden or neuritic dystrophy relative to the mean of the control (set at 100%). Immunization with each of the three N-terminal peptides significantly reduced amyloid burden (46–61%,  $P < 0.002$ ) (Fig. 2b). Furthermore, A $\beta$ 3–9 and A $\beta$ 5–11 significantly reduced neuritic pathology (34% and 41%, respectively;  $P < 0.05$ ) (Fig. 2c). In contrast, immunization with A $\beta$ 15–24 provided no protection against either amyloid burden or neuritic pathology. These results further support the association between plaque binding and antibody efficacy. They also indicate that capture of soluble A $\beta$  is not required for reduction of neuritic pathology, because the antibody response against A $\beta$ 3–9 provided strong plaque reactivity and the highest level of protection against neuronal dystrophy yet exhibited the weakest capacity for recognition of soluble peptide. These results, however, do not eliminate the possibility that antibodies specific for A $\beta$  capture could provide efficacy at higher titers or over longer periods of



**Fig. 1.** Antibodies produced by immunization with N-terminal fragments of A $\beta$  bind to amyloid plaques. (a) Peptides encompassing various domains of A $\beta$ 1–42 (synthesized contiguous to T cell epitope derived from ovalbumin) were used to immunize PDAPP mice. A reverse mer, A $\beta$ 5–1, was used as a negative control. (b) ELISA titers against aggregated A $\beta$ 1–42 were significantly higher over the length of the study in the A $\beta$ 5–11 and A $\beta$ 15–24 groups than in the A $\beta$ 1–5 group (1:14,457,  $P < 0.01$ , and 1:12,257,  $P < 0.05$  vs. 1:3,647, respectively; ANOVA followed by post hoc Tukey's test). (c) Unfixed cryostat sections from untreated PDAPP mouse brain were exposed to the sera of mice immunized with A $\beta$ 5–1, A $\beta$ 3–9, A $\beta$ 5–11, or A $\beta$ 15–24 (titers normalized to 1:1,000 for staining). Antibodies to A $\beta$ 15–24 did not bind to amyloid plaques. (Scale bar, 500  $\mu$ m.)

time, as has been reported by DeMattos *et al.* (6) using the high-affinity capture antibody 266.

**Antibody Isotype: IgG2a Antibodies Against A $\beta$  Are More Efficient than IgG1 or IgG2b Antibodies in Reducing Neuropathology.** Murine phagocytotic effector cells such as microglia within the CNS



**Fig. 2.** Capture of soluble A $\beta$ 1–42 by antibodies is not associated with reduced amyloid burden or neuritic pathology. (a) Sera from mice immunized with fragments of A $\beta$  were examined for their ability to capture radiolabeled soluble A $\beta$ 1–42 in an RIA. Sera from all animals immunized with A $\beta$ 15–24 were able to capture soluble A $\beta$ 1–42 (one serum sample had a titer higher than 1:1,350, and a precise titer was not determined) compared with 27% of those in the A $\beta$ 1–5 group and 3% of the A $\beta$ 3–9 group. Amyloid burden (b) and neuritic pathology (c) were evaluated using image analysis by a blinded microscopist. Values are expressed as a percentage of the mean of the A $\beta$ 5–1 group (negative control reverse mer peptide). The A $\beta$ 5–11 group was evaluated at a separate sitting from the other groups but in conjunction with the same negative control group as an internal reference (second A $\beta$ 5–1\* set, on the right). Amyloid burden was reduced significantly in the A $\beta$ 1–5, A $\beta$ 3–9, and A $\beta$ 5–11 groups ( $P < 0.001$ ). The bars represent median values, and the dashed horizontal line indicates the control level. Neuritic burden was reduced significantly in the A $\beta$ 3–9 and A $\beta$ 5–11 groups ( $P < 0.05$ ). Neither endpoint was altered significantly by immunization with the A $\beta$ 15–24 group. Statistical analysis was performed with square-root transformation (to normalize nonparametric distributions) and analyzed with ANOVA. A Dunnett's test then was used to compare the multiple groups A $\beta$ 1–5, A $\beta$ 3–9, and A $\beta$ 15–24 with their A $\beta$ 5–1 control and Mann-Whitney for the A $\beta$ 5–11 group with its corresponding A $\beta$ 5–1\* control.

express three different classes of IgG-specific Fc receptors (Fc $\gamma$  receptors): a high-affinity receptor, Fc $\gamma$ R1, and two low-affinity receptors, Fc $\gamma$ R2 and Fc $\gamma$ R3 (14). Fc $\gamma$ R2 is a single-chain re-

**Table 2. mAbs against A $\beta$ 3–7 have different avidity for aggregated and soluble synthetic A $\beta$ 1–42**

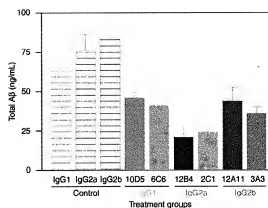
| Antibody | Epitope       | Isotype | ED <sub>50</sub> on aggregated A $\beta$ 1–42, pM | % Capture of soluble A $\beta$ 1–42 |
|----------|---------------|---------|---|-------------------------------------|
| 6C6      | A $\beta$ 3–7 | IgG1    | 40  | 1                                   |
| 10D5     | A $\beta$ 3–7 | IgG1    | 53  | 1                                   |
| 2C1      | A $\beta$ 3–7 | IgG2a   | 333   | 1                                   |
| 12B4     | A $\beta$ 3–7 | IgG2a   | 667   | 8                                   |
| 3A3      | A $\beta$ 3–7 | IgG2b   | 287   | 1                                   |
| 12A11    | A $\beta$ 3–7 | IgG2b   | 233   | 30                                  |

As a comparison, the antibody 266 at 10  $\mu$ g/ml would capture 70% of A $\beta$ 1–42.

ceptor with two major isoforms that apparently lack phagocytic capacity (15). Fc $\gamma$ R1 and Fc $\gamma$ R3 are heterooligomeric complexes in which the specific ligand-binding  $\alpha$  chains are associated with a common  $\gamma$  chain. The precise contribution of Fc $\gamma$ R1 and Fc $\gamma$ R3 to the phagocytosis of opsonized particles has not been defined; however, it has been shown that both receptors, and in particular Fc $\gamma$ R1, exhibit a higher affinity for murine IgG2a than for IgG1 or IgG2b (16). Furthermore, IgG2a has proven to be more effective in a number of *in vivo* clearance responses than the other antibody isotypes (17–20). Thus, if Fc-mediated phagocytosis of A $\beta$  peptide is an important mechanism for antibody-mediated plaque clearance, then IgG2a antibodies would be expected to reduce plaque burden more efficiently than the other antibody isotypes.

To address this issue, experiments were conducted with six mAbs: two of each IgG isotype and all directed against the same epitope of A $\beta$  (A $\beta$ 3–7). The epitope was defined further by amino acid substitution analysis; each antibody required the same three residues within A $\beta$ 3–7 for binding, and each could tolerate substitution within these residues by similar amino acids (data not shown). All the antibodies exhibited high avidity for aggregated A $\beta$ 1–42 ( $<1$  nM); however, the IgG1 antibodies showed  $\sim 10$ -fold greater binding avidity than the IgG2a antibodies ( $\sim 50$  vs.  $500$  pM) (Table 2). In contrast, only two of the antibodies could appreciably capture soluble A $\beta$ 1–42 at antibody concentrations of  $20 \mu$ g/ml: one of the IgG2b antibodies (12A11) and, to a lesser extent, one of the IgG2a antibodies (12B4). As a measure of their ability to trigger Fc-mediated plaque clearance, the six antibodies were compared in the *ex vivo* assay with primary mouse microglial cells and sections of brain tissue from PDAPP mice. Irrelevant IgG1, IgG2a, and IgG2b antibodies, having no reactivity toward A $\beta$  or other components of the assay, were used as isotype-matched negative controls. To quantify the degree of plaque clearance/A $\beta$  degradation that occurred by the end of the assay, A $\beta$  was extracted from the cultures of microglia and brain sections ( $n = 3$ ) with 8 M urea for analysis by ELISA (see Materials and Methods). As shown in Fig. 3, the two IgG2a antibodies against A $\beta$  reduced peptide levels in the cultures more efficiently (73% and 69%,  $P < 0.001$ ) than the IgG1 (28% and 35%, not significant) or IgG2b (48% and 59%,  $P < 0.05$  and  $0.001$ , respectively) antibodies. Because previous studies showed that *ex vivo* plaque clearance depends on Fc-receptor activity (7) and can occur in the presence of heat-inactivated serum, it is unlikely that complement played a significant role in mediating the effects observed in the *ex vivo* assay.

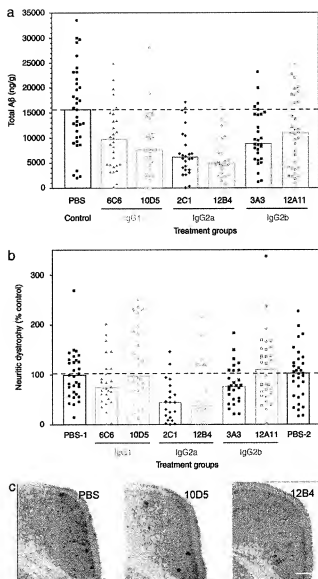
The antibodies then were investigated for *in vivo* efficacy. Antibody (10 mg/kg) or PBS control was administered by weekly i.p. injection for 6 months as described (7). Antibodies within all groups maintained similar serum titers against aggregated A $\beta$ 1–42 (1:3,500) with the exception of the IgG1 antibody 10D5,



**Fig. 3. Comparing antibodies against A $\beta$ 3–7 in the *ex vivo* assay.** IgG2a antibodies clear  $\beta$  amyloid plaque more efficiently than either the IgG1 or IgG2b isotypes. Murine primary microglial cells were cultured with unfixed cryostat sections of PDAPP mouse brain in the presence of antibodies of different isotypes directed against A $\beta$ 3–7. Irrelevant IgG1, IgG2a, and IgG2b antibodies were used as the respective isotype-matched negative controls. After 24 h of incubation, the total level of A $\beta$  remaining in the cultures was measured by ELISA. The two anti-A $\beta$  IgG2a antibodies reduced A $\beta$  levels in the cultures (69% for 2C1 and 73% for 12B4;  $P < 0.001$ ) more efficiently than the IgG2b isotype antibodies (48% for 12A11,  $P < 0.05$ , and 59% for 3A3,  $P < 0.001$ ). The anti-A $\beta$  IgG1 antibodies did not significantly reduce A $\beta$  levels. Data were analyzed with ANOVA followed by a post hoc Dunnett's test.

which displayed 3- to 4-fold higher titers ( $\sim 1:13,000$ ). At the end of the study, total levels of cortical A $\beta$  were determined by ELISA. Although each of the antibodies significantly reduced total A $\beta$  levels compared with the PBS control ( $P < 0.001$ ) (Fig. 4a), there was a trend toward greater levels of reduction by the two IgG2a antibodies (61% and 69% reduction) than by the IgG1 (38% and 52%) or the IgG2b (44% and 31%) antibodies. These results suggest that the higher affinity of IgG2a for Fc receptors is an important parameter for clearance. Furthermore, because murine IgG1 antibodies cannot fix complement but provided levels of A $\beta$  clearance comparable to the complement-fixing IgG2b antibodies, complement does not seem to play a critical role during the clearance process *in vivo*.

The level of neuritic dystrophy then was examined in sections of brain tissue from the mice to determine the association between plaque clearance and neuronal protection. Again, data are shown for individual animals and expressed as the percentage of neuritic dystrophy relative to the mean of the control (set at 100%). Although all the antibodies triggered plaque clearance, only the IgG2a antibodies provided significant reduction in neuritic dystrophy (12B4,  $P < 0.05$ , and 2C1,  $P < 0.001$ ) (Fig. 4b and c). Interestingly, the antibody 10D5 (IgG1) was less effective than either of the IgG2a antibodies even though it exhibited higher avidity for aggregated A $\beta$ 1–42 (Table 2) as well as amyloid plaques (data not shown) and maintained significantly higher serum titers than the other antibodies. Thus, antibody isotype and affinity for Fc receptors seem to be important attributes for both clearance of A $\beta$  and protection against neuritic dystrophy and may be more important than the relative avidity of antibodies for A $\beta$ 1–42 (Table 2). Also, these studies confirmed the observation obtained in the peptide immunization study described above that antibodies do not need to strongly capture soluble A $\beta$ 1–42 to provide protection against neuritic dystrophy. The antibody 12A11 (IgG2b) captured soluble monomeric A $\beta$ 1–42 more efficiently than either of the IgG2a antibodies (Table 1) but was not as effective. Also, both IgG2a antibodies provided similar protection even though only one could capture the soluble peptide detectably. The results of the current study are not necessarily inconsistent with other



**Fig. 4.** Anti- $A\beta$  IgG2a antibodies reduced AD-like neuropathology more efficiently than other isotypes *in vivo*. PDAPP mice received weekly i.p. injections of antibodies starting at 12 months of age for 6 months. (a) Total  $A\beta$  levels shown for individual mice sorted by treatment group ( $n = 30$ ). The bars represent median values, and the dashed horizontal line indicates the control level. Although  $A\beta$  levels were reduced significantly in all antibody groups ( $P < 0.001$  vs. PBS; ANOVA followed by post hoc Dunnett's test), the IgG2a groups exhibited the highest degree of clearance. (b) The percentage of frontal cortex occupied by neuritic dystrophy was determined by image analysis. The different groups within this experiment were analyzed in two sets by using the same PBS group as an internal standard (PBS-1 and PBS-2). PBS-1 was the control for the 6C6, 10D5, 12B4, and 12A11 groups, and PBS-2 was the control for the 2C1 and 3A3 groups. To compare the groups, values for individual animals are expressed as a percentage of the mean of their respective PBS control group (set at 100%). The bars represent median values, and the dashed horizontal line indicates the control level. Neuritic dystrophy was reduced significantly only by the IgG2a isotype antibodies (12B4,  $P < 0.05$ , and 2C1,  $P < 0.001$ ; ANOVA followed by post hoc Dunnett's test). (c) Dystrophic neurites were labeled with the amyloid precursor protein-specific antibody 8E5, and were found in association with plaques. Relative to PBS control, the neuritic pathology was reduced significantly in animals treated with 12B4 but not 10D5. (Scale bar, 250  $\mu$ m).

investigations and hypotheses surrounding anti- $A\beta$ -based immunotherapies. Duration of treatment, route of administration, and specific antibody properties all likely have important effects on

the observed outcomes. For example, Solomon *et al.* (5) suggested that anti- $A\beta$  antibodies may directly inhibit or reverse amyloid fibril formation. Bacsakai *et al.* (8) provided support for the possible direct dissolution mode of action by showing removal of plaque *in vivo* by F(ab')<sub>2</sub> fragments of an anti- $A\beta$  antibody after direct application to the brain. It should be noted, however, that *in vitro* dissolution was reported to be restricted to antibodies against  $A\beta$ 3–6 (21), whereas the antibody used in the *in vivo* study was against  $A\beta$ 1–5. In addition, we show that antibodies against fragments  $A\beta$ 1–5,  $A\beta$ 3–9, and  $A\beta$ 5–11 are all capable of reducing plaque burden. Thus the mechanism of plaque reduction does not seem to have the same restricted epitope as reported for *in vitro* fibril dissolution. Although the direct application of high-dose antibody to brain was capable of clearing plaque without Fc-mediated phagocytosis, the present data demonstrate that the efficiency in reducing plaque burden and neuritic dystrophy is best when the antibody isotype maximally supports phagocytosis and that efficacy can be achieved by antibodies of several epitope specificities that all are capable of binding plaque *in vivo* or *ex vivo*.

Another reported mechanism of efficacy is through capture of soluble  $A\beta$  (6). The use of antibody 266, a high-affinity capture antibody, at concentrations sufficient to produce detectable cerebrospinal fluid levels reduced plaque burden after chronic treatment. It is likely that the capture titer achieved by immunization with the  $A\beta$ 15–24 fragment, although still substantially greater than the other immunogens tested in this study, is less than that achieved by 266 dosing, and this may impact the outcome. Thus it may be possible to achieve the same endpoint through multiple mechanisms including a chronic capture of  $A\beta$  species, dissolution of plaques, or phagocytosis of existing aggregates.

In summary, although antibodies against  $A\beta$  may exhibit efficacy in a number of ways, protection against AD-like neuropathology can be obtained by antibodies that bind to plaques and trigger Fc-mediated clearance. IgG2a antibodies, which exhibit higher affinity than other isotypes for phagocytic Fc receptors (in particular FcγR1), provided the highest level of plaque clearance and were the only anti- $A\beta$  antibodies to provide neuronal protection under the conditions tested. Plaque clearance seemed independent of complement activation, because IgG1 antibodies, which cannot fix complement, were as effective as the complement-fixing IgG2b antibodies. These results are consistent with the role of high-affinity Fc receptors in other clearance systems, where they have been shown to be particularly effective for inducing clearance in conditions with low antibody concentrations (as would be anticipated in the CNS). The density of target-bound IgG that is required for complement activation has been reported to be higher than that required for Fc receptor-mediated phagocytosis (17). Accordingly, complement involvement in plaque clearance may be more pronounced at higher doses of antibody, where there would be an increased density of antibody bound to plaques. Interestingly, in contrast to other macrophage-activation paradigms, it has been shown that activation of phagocytic cells through Fc receptors results in production of the antiinflammatory cytokine IL-10 and inhibition of proinflammatory IL-12 (22). Thus, antibodies against  $A\beta$  may allow resolution of an otherwise chronic, unresolved inflammatory response associated with plaques in AD by both clearing  $A\beta$  and altering the inflammatory environment.

In addition, the current studies demonstrate that antibody epitopes within the N terminus of  $A\beta$  are important for plaque clearance and neuronal protection via an Fc-mediated mechanism. Passive administration of mAbs against defined  $A\beta$  epitopes reduced plaque burden and neuritic pathology to the same degree as active immunization. Although the 5- to 7-aa residue epitopes used for immunization in our studies are

themselves too short to elicit T cell help for antibody production, we showed that these epitopes can be synthesized in conjunction with an exogenous T cell epitope to produce an efficacious antibody response after administration. Such an approach will avoid generation of T cell immunity against A $\beta$  as a self-antigen and may preclude the potential issues with encephalitis that were observed recently with a subset of patients in the clinic after immunization with whole A $\beta$ 1–42 (23, 24). Likewise, T cell

immunity will not be elicited with passive administration of antibodies against A $\beta$ . Thus, both immunoconjugates containing defined epitopes of A $\beta$  and mAbs against appropriate A $\beta$  epitopes offer excellent alternatives to whole-peptide immunization for the treatment of AD.

We thank Chuck Davies for advice in the statistical analysis and Dr. Manuel Buttini for helpful editorial comments.

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